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(54) **Closure device for transcatheter operation and catheter assembly therefor**

Verschlussvorrichtung für Kathetereingriff, sowie Kathetereinrichtung dafür

Prothèse de fermeture à mettre en place par l'intermédiaire d'un cathéter

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(56) References cited:
WO-A-98/27868 **US-A- 5 108 420**
US-A- 5 829 447

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Description

[0001] The present invention relates to a closure device and a catheter assembly suitable for transcatheter operations, i.e., operations for closing or repairing endocardiac or vascular defects.

[0002] At present, the Fontan operations have been performed as the repair operations of complex cardiac anomalies such as a single ventricle. In this operation, the cardiac function is repaired by employing a single ventricle for the systemic circulation system, and by directly leading the blood from the vein of the systemic circulation system to the pulmonary artery in the pulmonary circulation system. After this operation, however, extreme decrease in cardiac output may occur because of transient ventricular hypofunction or increase in pulmonary vascular resistance, associated with vasospasm or residual peripheral pulmonary arteriostenosis. Thus, it occasionally becomes difficult to perform postoperative management.

[0003] In order to avoid increase of the pulmonary vascular resistance and decrease of the cardiac output, therefore, it is general practice for the operation to form a small fenestration or perforation of about 4 mm between atria (in most cases, the interatrial septum is made of an artificial membrane) to allow the blood to flow through the small fenestration or perforation. This procedure followed by formation of the small fenestration or perforation is referred to as the Fontan fenestration. The surgical operation is generally carried out as a method of treatment for closing the small fenestration after the Fontan fenestration. As a matter of course, however, this surgical operation is followed by thoracotomy of a patient in addition to the treatment of the affected area, which imposes a burden on the patient. In particular, in case of child patients, this burden is considerably large.

[0004] Percutaneous transluminal therapeutic catheterization is known as a noninvasive procedure for repairing endocardial defects. In this procedure, an atrial septal defect is closed by transvenously inserting an intercardiac catheter into the heart, and closing the atrial septal defect with an occlusion device. The first clinical success of this method was reported by King and Mill in 1976. In the method carried out by King et al, closure of the atrial-septal defect is carried out by introducing left and right atrial double umbrellas into an atrium with an insertion tool composed of a core wire and a double-layered catheter coaxially arranged thereon, fitting the umbrellas on the interatrial septum so that the interatrial septum is sandwiched between them, and then combining the umbrellas into one.

[0005] However, this method can not be applied to children, especially, to preschool children since it uses a very thick insertion tool and hard umbrellas. To solve this problem, Rashkind investigated miniaturization of the device, developed in 1977 a single umbrella type occlusion plug with hooks, and reported its clinical success in children. However, this procedure has another problem such that the plug may be occasionally caught in an unintended site of the heart because of its hooks. Thus, when once the umbrella was opened in the heart, the umbrella-like member can not be changed its position or withdrawn from the site, resulting in emergency operation. To overcome such disadvantages, Rashkind further developed an improved occlusion plug comprising two umbrellas united one another. This plug has widely been put into clinical use for closure of patent ductus arteriosus.

[0006] On the other hand, Lock et al developed a clamshell-shaped intraatrial occlusion device and applied for a patent (JP-A 5-237128), which has been modified by additionally providing coil springs for the double umbrella occlusion plug of Rashkind so that a thin interatrial septum is firmly sandwiched between two umbrellas under the action of the coil springs arranged at each middle portion of eight stainless steel frames of the plug.

This occlusion device is so designed as to be firmly fitted on the thin interatrial septum when the two umbrellas members are in close contact with one another under the overlapped conditions. Because of its configuration similar to a clamshell of a bivalve, this device is called a "clamshell septal occluder". In use, the treatment is carried out by introducing an elongated sheath of 11 French thick into the repair site through a femoral vein of a patient. Since this method is applicable to a patient with a weight of more than 8 kg, it has been widely used as a percutaneous transluminal therapeutic transcatheter closure of atrial septal defects.

[0007] The closure treatment of the small hole after Fontan fenestration is done with the occlusion plug by percutaneous transluminal therapeutic transcatheterization.

[0008] However, in the method of closing the small hole caused by the Fontan fenestration with the occlusion plug of the prior art, adhesion of thrombus may occur because of a large occlusion plate of the plug, resulting in high risk of complications.

[0009] Further, if disembarrassment or dislodgement of the occlusion device has taken place, it is difficult to withdraw the occlusion plug because of its shape or size.

[0010] In US-A-5 108 420 an aperture occlusion device is composed of two biocompatible membranes, wherein wire ribs are secured to each of the biocompatible membrane. The wire ribs are made of a shape memory retentive material. Further an eye is attached to the wire ribs which extends from the center of the aperture occlusion device and engages a knuckle of a release wire.

[0011] The present invention has been made in view of the above-mentioned circumstances and aimed at providing a closure device for transcatheter operations, that is simple in shape, easy to operate, capable of closing a small hole safely and infallibly, easy to withdraw and less in adhesion of thrombus. Also, the present invention is aimed at providing

a catheter assembly for introducing the closure device into the site of operation.

[0012] As the result of assiduous studies for solution of the above problem, the present inventors have conceived an idea of providing a closure membrane with a fixing means of fixing the closure membrane to a tissue wall surrounding a defect aperture, said fixing means exhibiting a shape memory behavior and recovering its memorized shape when heated to the body temperature to sandwich the tissue wall from both sides of the defect aperture between the fixing means and the closure membrane.

[0013] In other words, a subject matter of the present invention is directed to an easily foldable closure device for transcatheter operations, comprising: a flat closure member including a closure membrane and a frame at the periphery of the membrane; a fixing member for fixing the closure member to a tissue surrounding a defect aperture; and a connecting member fixed at one end to the fixing member and at the other end to the frame of said closure member for holding them in close proximity one another. The frame, fixing member and connecting member are respectively made of the same or different shape-memory alloys with a shape recovery temperature of 30 to 37 °C.

[0014] According to the present invention, there is provided a closure device for transcatheter operations, comprising: a flat closure member including a closure membrane made of a fabric or non-woven fabric of a bio-compatible material, and a fringing frame at the periphery of said closure membrane, said fringing frame being made of a shape-memory alloy with a shape recovery temperature of 30 to 37 °C; a fixing member for fixing said closure member to a tissue surrounding a defect aperture, said fixing member being approximately parallel to one plane of said closure member and adapted to be located on one side of said defect aperture so that said tissue is sandwiched between said fixing member and said closure member located on the opposite side of said defect aperture, said fixing member being made of a shape-memory alloy with a shape recovery temperature of 30 to 37 °C; and a connecting member fixed at one end to said fixing member and at the other end to the above-mentioned frame of said closure member for holding said closure member and fixing member in close proximity one another, said connecting member being made of a shape-memory alloy with a shape recovery temperature of 30 to 37 °C, said closure device being easily foldable at a temperature lower than the shape recovery temperature of said shape memory alloys.

[0015] In an embodiment of the present invention, the fringing frame, the fixing member and the connecting member may be formed in one united body. In this case, it is preferred to cover a joint portion of the connecting member and the fringing frame with a tubular covering member to prevent the fixing member from being reversed.

[0016] Further, a base of the fixing member may be stitched to the closure membrane of the closure member with a thread so that the defect aperture is located in the central portion of the closure member when the defect aperture is closed.

[0017] In addition, the closure member may be provided with a grappling hole to make it easy to hold the closure device by the holding means of the catheter assembly.

[0018] According to the invention, there is also provided a catheter assembly for use in combination with the occlusion device defined as above, comprising a sheath opened at both ends and being able to accommodate a folded closure device of claim 1; and, an operating rod provided at a distal end thereof with a holding means for releasably holding said closure device, said operating rod being insertable into said sheath from one end thereof under conditions of holding said closure device and being able to unsheath said closure device through the opposite end of said sheath.

[0019] In an embodiment of the present invention, it is preferred to constitute the holding means with an easily flexible linear member extending from the distal end of the operating rod in the axial direction thereof.

[0020] The embodiment of the present invention will be explained below on the basis of the accompanying drawings.

Fig. 1 is a plane view illustrating one embodiment of a closure device of the present invention;

Fig. 2 is a plane view illustrating another embodiment of the closure device of the present invention;

Fig. 3 is a plane view illustrating still another embodiment of the closure device of the present invention.

Fig. 4 is a vertical section view illustrating one embodiment of a catheter assembly of the present invention.

Fig. 5 is a view illustrating the closure device of Fig. 1 put in the catheter assembly shown in Fig. 4.

Fig. 6 is an illustration of a percutaneous closure treatment employing the closure device and catheter assembly of the present invention.

Fig. 7 is an illustration of a percutaneous closure treatment employing the closure device and catheter assembly of the present invention.

Fig. 8 is an illustration of a percutaneous closure treatment employing the closure device and catheter assembly of the present invention.

[0021] Firstly, the explanation will be made on the closure device [C] for transcatheter operations of the present invention.

[0022] As shown in Figs. 1 to 3, the closure device C of the present invention comprises an easily foldable closure member 1, a fixing member 2 for fixing the closure member 1 to a tissue surrounding a defect aperture, and a connecting member 3 for holding the closure member 1 and the fixing member 2 in close proximity one another. The connecting member 3 is fixed at one end to the fixing member 2 and at the other end to the frame 12 of said closure member 1.

[0023] The closure member 1 is composed of a closure membrane 11 made of a bio-compatible material and a fringing frame 12 made of a shape-memory alloy with a shape recovery temperature of 30 to 37 °C at the peripheral portion of the closure membrane 11. The closure member 1, the fixing member 2 and the connecting member 3 may be formed in one united body. Also, the closure membrane 11 of the closure member 1 may be fixed to the base 21 of the fixing member 2 with a thread 5 passing through the central portion of the closure membrane 11 of the closure member 1 so that the defect aperture D is located at the central portion of the closure member 1 when the defect aperture D is closed.

[0024] The closure member 1 is an easily foldable flat member and is preferably composed of the closure membrane 11 made of fabrics or nonwoven fabrics of bio-compatible materials, and the frame 12 made of a shape-memory alloy with a shape recovery temperature of 30 to 37 °C and arranged on the peripheral portion of the closure membrane 11. Although the closure member 1 may take any configuration, it is preferred for the closure member 1 to have a flat configuration which is easy to fold and analogous to a shape of the defect aperture D (generally, in a circle), such as a circular, hexagon, octagon and the like. However, the closure member 1 is not limited to these shapes. It is sufficient for the closure member 1 to have a size large enough to close the defect aperture D. In order to assure the closure of the defect aperture D, it is preferred to use the closure member 1 with a size of 1.5 to 2 times the defect aperture D.

[0025] Further, the closure member 1 may be provided with a hole 4 to make it easy to hold the closure member with the holding means of the catheter assembly mentioned below.

[0026] As the biocompatible material for closure membrane 11, there may be used polyesters such as polyethylene terephthalate; polyolefins such as polyethylene and polypropylene; polyamides such as nylon 6 and nylon 66; fluoroplastics such as polytetrafluoroethylene and polyvinylidene fluoride, chlorine-containing resins such as polyvinyl chloride and polyvinylidene chloride; polyurethane; semisynthetic resins such as cellulose plastics; natural fibers; or the like. These materials may be used in the form of fabrics, nonwoven fabrics, films, porous sheets, or their composites.

[0027] The frame 12 is made of a shape-memory alloy having a shape recovery temperature of 30 to 37 °C in the form of a line or wire. Preferred shape-memory alloys generally include Ni-Ti alloys, Cu-Zn-Al alloys, Cu-Al-Ni alloys and the like.

The fixing member 2 for fixing the closure member 1 to the tissue surrounding the defect aperture D is a member to be located on one side of the tissue surrounding the defect aperture D to sandwich the tissue between the fixing member 2 and the closure member 1 located on the opposite side of the tissue surrounding the defect aperture D. The fixing member 2 is made of a shape-memory alloy similar to that of the frame 12 in the form of an estoile composed of a base 21 and plural arms 22 (generally, two to four elements) extending radially from the base 21 in one plane. The fixing member 2 is fixed to the connecting member 3 so that the arms 22 are approximately parallel to the plane of the closure member 1. The connecting member 3 may be united with the fixing member 2 by suitable connecting means such as welding. In this case, the connected portion between the connecting member 3 and the fixing member 2 is preferably covered with and fixed by a covering member 31 so as to prevent the fixing member 2 from reversing its course.

[0028] The connecting member 3 is generally made of the same shape-memory alloy as that used for the frame 11 and fixing member 2 in the form of a slender linear member, and fixed at its one end to the closure member 1 and at the opposite end to the fixing member 2 to hold them in close proximity one another. In order to hold the fixing member 2 substantially parallel to the plane of the closure member 1, the connecting member 3 is arranged in approximately parallel to the plane of the closure member 1.

[0029] In use, the connecting member 3 is folded first together with the closure member 1 and the fixing member 2 folded in the longitudinal axis of a sheath 6 mentioned below, and then inserted into the sheath 6 of a catheter assembly A shown in Fig. 4. During passage of the closure device through the sheath 6, the frame 12, fixing member 2 and connecting member 3 are heated to their recovery temperature by the blood, but they are kept in the folded condition by the sheath as shown in Fig. 5. When the frame 12, arms 22 and connecting member 3 are pushed out of the sheath 6, they are allowed to recover respective original shapes so that the tissue surrounding the defect aperture D is sandwiched between the closure member 1 and fixing member 2.

Next, explanation will be made on the catheter assembly A of the present invention.

[0030] As shown in Fig. 4, the catheter assembly A of the present invention comprises a sheath 6 and an operating rod 7 for introducing the aforesaid closure device C into the operation site through the sheath 6 to close the defect aperture D with the closure device.

[0031] The sheath 6 is a tubular member opened at both ends thereof and has a lumen 61 for holding the folded closure device C therein and for movably holding the operating rod 7. The sheath 6 is provided at its proximal end with a connector 62 having a port 621 for insertion of the operating rod 7, and a stepped through-hole composed of a small-sized portion 623 and a large-sized portion 624 provided coaxially with the small-sized portion 623.

[0032] The connector 62 is provided with hemostatic means (e.g., a hemostatic valve) for preventing the blood from leakage during the operations, and a lateral tube 64 for infusion of heparinized physiological saline to prevent coagulation of blood during the operations. The hemostatic means used in this embodiment is composed of a gasket or packing 63 having a through-hole provided in a central part thereof. The packing 63 is set in the large-sized portion 624 and forced against stepped portion 625 of the connector 62 by a bolting member 622 having a through-hole at the central portion

thereof.

[0033] As a material for the sheath 6, there may be used any one of the materials selected from the group consisting of fluoroplastics such as polytetrafluoroethylene; synthetic resins such as polypropylene and polyethylene, polyesters, polyurethane; and meshed or coiled stainless steels (e.g., SUS 304).

[0034] The operating rod 7 is comprised of a rod portion 71, a holding means 72 provided at a distal end of the rod portion 71 for releasably holding the closure device C, and a handle 73 for manipulating the operating rod 7 by hand. By manipulating the operating rod, the folded closure device C held by the holding means 72 is inserted into the sheath 6 and then pushed out of the opposite end of the sheath 6.

[0035] The holding means 72 is preferably made of an easily flexible linear member extending in the axial direction of the operating rod 7. In this case, a material for the rod portion 71 may be the same material as those used for the sheath 6. As a material for the holding means 72, it is preferred to use an easily flexible elastic material such as, for example, metals such as stainless steels and brasses; flexible plastics such as polypropylene and polyester; super elastic alloys and the like. Preferably, as shown in Fig. 4, the holding means 72 is turned at its proximal portion 721 to form an annular portion that makes it easy to inflect the holding means.

[0036] The holding means 72 is turned to hold the closure device C and then inserted into the sheath 6. When the closure device C is pushed out of the sheath 6 by operating the operating rod 7, the holding means 72 is returned to its original shape extending in the axial direction of the operating rod 7 since the holding means 72 is made of an elastic material. Thus, the closure device C is released from the holding means 72.

[0037] Next, the use of the catheter assembly A of the present invention will be explained making reference to Figs. 5 to 8 which show a series of procedures in percutaneous closure treatment employing the closure device and catheter assembly of the present invention.

[0038] First of all, the holding means 72 of the operating rod 7 is passed through a part of the closure membrane 11 adjacent to the frame 12 of the closure device C. Then, the closure device C is folded as shown in Fig. 5 (the fixing member 2 and the connecting member 3 are inflected in the longitudinal direction) and then insert into the sheath 6. In this case, the holding means 72 is inserted into the sheath 6 under the condition of being inflected in the direction of the handle 73 to hold the frame 12 of the closure device C.

[0039] Next, the catheter assembly A is inserted into an elongated sheath (not illustrated in the figures) previously introduced in a right atrium through a femoral vein of a right leg, to introduce the distal end of the sheath 6 into the right atrium. After the distal end of the catheter assembly A is protruded from the defect aperture D of the interatrial septum into the left atrium through, the fixing member 2 of the closure device C is pushed out into the left atrium by pushing the operating rod 7. Simultaneously therewith, the fixing member 2 recovers its shape and returns to its original state parallel to the closure member 1, so that the fixing member 2 becomes impossible to pass through the defect aperture D (Fig. 6).

[0040] The catheter assembly A is then pulled backward so that its distal end is located in the right atrium. By further pushing the operating rod 7 into the sheath 6, the closure member 1 of the closure device C is pushed out of the sheath 6 in the right atrium and then recovers its unfolded original shape (cf. Fig. 7). The operating rod 7 is further pushed into the sheath 6 (if necessary, with the sheath 6 being pulled back.) to protrude the holding means 72 from the sheath 6. As soon as the holding means 72 is wholly protruded from the sheath 6, it returns to its original shape, i.e., the shape extending in the axial direction of the rod portion 71, thereby disengaging the closure device C. Simultaneously therewith, the recovered closure member 1 is pulled toward the fixing member 2 by elasticity of the frame 12 and connecting member 3 so that the tissue surrounding the defect aperture D is sandwiched between the closure member 1 and the fixing member 2 to close the defect aperture D (cf. Fig. 8). Thus, the operation is finished.

[0041] As will be understood from the above description, a defect aperture is closed easily and infallibly through the closure device and catheter assembly for transcatheter operations of the present invention. In addition, the adhesion of thrombus to a closure device, the problem with the use of a closure device of the prior art, can be kept down to minimum therethrough. Also, the closure device can be easily withdrawn from the site even if disembarassment or dislodgement of the closure device had taken place.

Claims

1. A foldable closure device for transcatheter operations, comprising:

a closure member (1), including a closure membrane (11) made of a fabric or a non-woven fabric of a bio-compatible material, and a frame (12) at the periphery of said closure membrane, said frame being made of a shape-memory alloy with a shape recovery temperature of 30 to 37 °C;

a fixing member (2) for fixing said closure member (1) to a tissue surrounding a defect aperture (D), said fixing member (2) being approximately parallel to one plane of said closure member (1) and adapted to be located on one side of said defect aperture (D) so that said tissue is sandwiched between said fixing member (2) and

said closure member (1) located on the opposite side of said defect aperture, said fixing member being made of a shape-memory alloy with a shape recovery temperature of 30 to 37 °C; and
a connecting member (3) fixed at one end to said fixing member (2) and at the other end to the above-mentioned frame of said closure member (1) for holding said closure member (1) and fixing member (2) in close proximity to one another, said connecting member (3) being made of a shape-memory alloy with a shape recovery temperature of 30 to 37 °C.

2. The closure device according to Claim 1, wherein the frame (12) of the closure member (1) is united with said fixing member (2) and said connecting member (3).
3. The closure device according to claim 2, wherein a connected portion of said fixing member (2) and said connecting member (1) is covered with a tubular covering member (31).
4. The closure device according to any one of claims 1 to 3, wherein a base of the fixing member (2) is fixed to the closure membrane (11) of the closure member (1) with a thread (4).
5. The closure device according to any one of claims 1 to 4, wherein the closure member (1) is provided with a hole.
6. A catheter assembly comprising;
a sheath (6) having openings at both ends
an operating rod (7) provided at a distal end thereof with a holding means (72) for releasably holding said closure device, said operating rod (7) being insertable into said sheath (6) from one end thereof under conditions of holding said closure device and being able to unsheath the closure device through the opposite end of said sheath (6), and a folded closure device of any one of claims 1 to 5.
7. The catheter assembly according to claim 6, wherein said holding means (72) is made of an easily flexible linear member extending from the distal end of the operating rod in the axial direction thereof.

Patentansprüche

1. Faltbare Verschlussvorrichtung für Transkathetereingriffe mit:

einem Verschlussstück (1) mit einer Verschlussmembran (11) aus einem Gewebe oder einem Textilverbundstoff (oder Vliesstoff) aus einem biokompatiblen Material und einem Rahmen (12) am Umfang der Verschlussmembran, wobei der Rahmen aus einer Formgedächtnis-Legierung mit einer Formgedächtnis-Temperatur von 30 bis 37°C hergestellt ist;
einem Fixierteil (2) zum Fixieren des Verschlussstücks (1) an einem Gewebe, das eine Defekt-Apertur (D) umgibt, wobei das Fixierteil (2) annähernd parallel zu einer Ebene des Verschlussstücks (1) ist und auf einer Seite der Defekt-Apertur (D) angeordnet werden kann, so daß das Gewebe zwischen dem Fixierteil (2) und dem Verschlussstück (1), das sich auf der gegenüberliegenden Seite der Defekt-Apertur befindet, sandwichartig angeordnet ist, wobei das Fixierteil aus einer Formgedächtnis-Legierung mit einer Formgedächtnis-Temperatur von 30 bis 37 °C hergestellt ist; und
einem Verbindungsteil (3), das mit einem Ende an dem Fixierteil (2) und mit dem anderen Ende an dem vorstehend erwähnten Rahmen des Verschlussstücks (1) fixiert ist, um das Verschlussstück (1) und das Fixierteil (2) in enger Nachbarschaft zueinander zu halten, wobei das Verbindungsteil (3) aus einer Formgedächtnis-Legierung mit einer Formgedächtnis-Temperatur von 30 bis 37 °C hergestellt ist.

2. Verschlussvorrichtung nach Anspruch 1, wobei der Rahmen (12) des Verschlussstücks (1) mit dem Fixierteil (2) und dem Verbindungsteil (3) vereinigt ist.
3. Verschlussvorrichtung nach Anspruch 2, wobei ein verbundener Abschnitt des Fixierteils (2) und des Verbindungsteils (1) mit einem röhrenförmigen Hüllteil (31) umhüllt ist.
4. Verschlussvorrichtung nach einem der Ansprüche 1 bis 3, wobei eine Basis des Fixierteils (2) an der Verschlussmembran (11) des Verschlussstücks (1) mit einem Faden (4) fixiert ist.

5. Verschlussvorrichtung nach einem der Ansprüche 1 bis 4, wobei das Verschlussstück (1) mit einem Loch versehen ist.

6. Katheteranordnung mit:

5 einem Schaft (6) mit Öffnungen an beiden Enden;
einem Führungsstab oder -draht (7), der an seinem distalen Ende mit einer Halteeinrichtung (72) zum lösbaren Halten der Verschlussvorrichtung versehen ist, wobei der Führungsstab (7) von seinem einen Ende in den Schaft (6) unter den Bedingungen einführbar ist, daß die Verschlussvorrichtung gehalten wird, und die Verschlussvorrichtung durch das entgegengesetzte Ende des Schafts (6) hinausgeschoben werden kann, und
10 einer gefalteten Verschlussvorrichtung nach einem der Ansprüche 1 bis 5.

7. Katheteranordnung nach Anspruch 6, wobei die Halteeinrichtung (72) aus einem leicht flexiblen geraden Teil besteht, das sich vom distalen Ende des Führungsstabs in seiner axialen Richtung erstreckt.

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Revendications

1. Dispositif de fermeture pliable pour les opérations de trans-cathétisme, comprenant:

20 un élément de fermeture (1) comprenant une membrane de fermeture (11) réalisée avec un tissu ou un tissu non tissé d'un matériau biocompatible et un châssis (12) au niveau de la périphérie de ladite membrane de fermeture, ledit châssis étant réalisé avec un alliage à mémoire de forme avec une température de recouvrement de forme de 30 à 37°C ;
un élément de fixation (2) pour fixer ledit élément de fermeture (1) sur un tissu entourant une ouverture de défaut (D), ledit élément de fixation (2) étant approximativement parallèle à un plan dudit élément de fermeture (1) et adapté pour être situé d'un côté de ladite ouverture de défaut (D), de sorte que ledit tissu est pris en sandwich entre ledit élément de fixation (2) et ledit élément de fermeture (1) situé du côté opposé de ladite ouverture de défaut, ledit élément de fixation étant réalisé avec un alliage à mémoire de forme avec une température de recouvrement de forme de 30 à 37°C ; et
30 un élément de raccordement (3) fixé au niveau d'une extrémité sur ledit élément de fixation (2) et au niveau de l'autre extrémité sur le châssis mentionné ci-dessus dudit élément de fermeture (1) pour maintenir ledit élément de fermeture (1) et ledit élément de fixation (2) en étroite proximité l'un de l'autre, ledit élément de raccordement (3) étant réalisé avec un alliage à mémoire de forme avec une température de recouvrement de forme de 30 à 37°C.

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2. Dispositif de fermeture selon la revendication 1, dans lequel le châssis (12) de l'élément de fermeture (1) est assemblé audit élément de fixation (2) et audit élément de raccordement (3).

3. Dispositif de fermeture selon la revendication 2, dans lequel une partie raccordée dudit élément de fixation (2) et dudit élément de raccordement (1) est recouverte avec un élément de recouvrement tubulaire (31).

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4. Dispositif de fermeture selon l'une quelconque des revendications 1 à 3, dans lequel une base de l'élément de fixation (2) est fixée sur la membrane de fermeture (11) de l'élément de fermeture (1) avec un filetage (4).

45 5. Dispositif de fermeture selon l'une quelconque des revendication 1 à 4, dans lequel l'élément de fermeture (1) est prévu avec un trou.

6. Ensemble de cathéter comprenant :

50 une gaine (6) ayant une ouverture (5) aux deux extrémités ;
une tige de commande (7) prévue au niveau de son extrémité distale avec des moyens de support (72) pour supporter de manière amovible ledit dispositif de fermeture, ladite tige de commande (7) étant insérable dans ladite gaine (6) depuis son extrémité dans des conditions de support dudit dispositif de fermeture et pouvant faire sortir le dispositif de fermeture de la gaine à travers l'extrémité opposée de ladite gaine (6) et un dispositif de fermeture plié selon l'une quelconque des revendications 1 à 5.

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7. Ensemble de cathéter selon la revendication 6, dans lequel lesdits moyens de support (72) sont réalisés avec un élément linéaire facilement flexible s'étendant à partir de l'extrémité distale de la tige de commande dans sa direction axiale.

Fig. 1

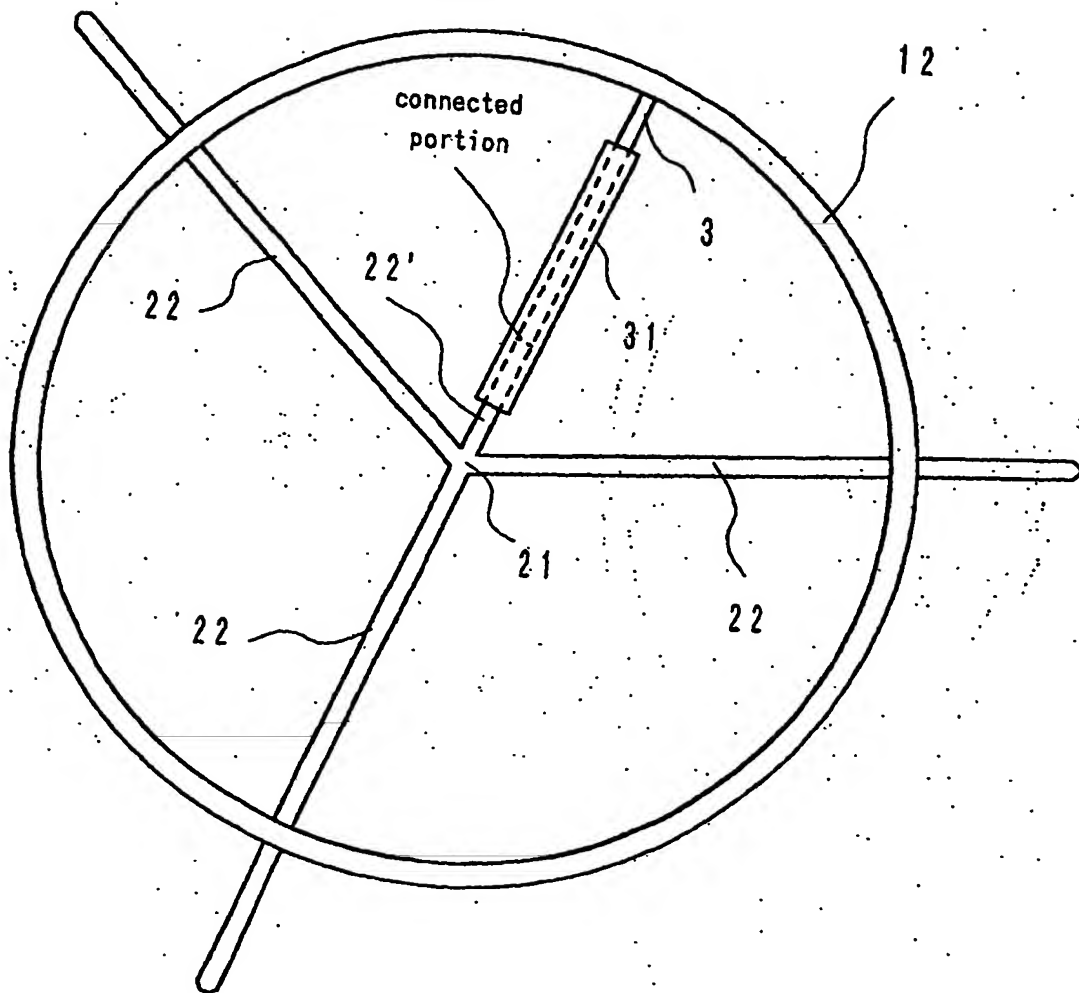


Fig. 2

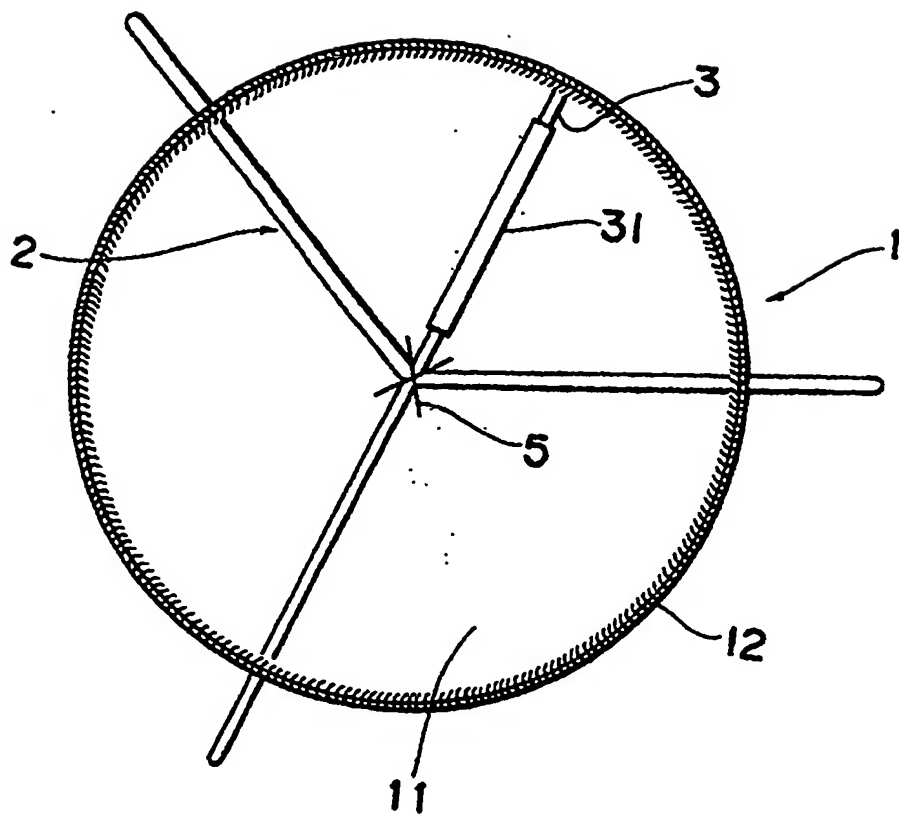


Fig. 3

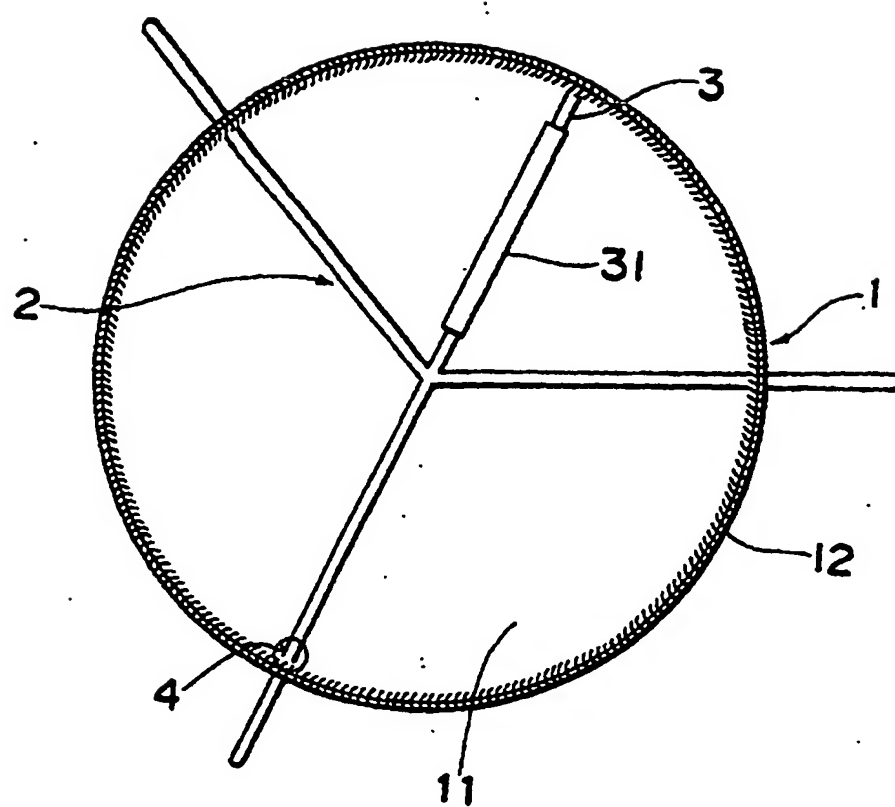


Fig. 4

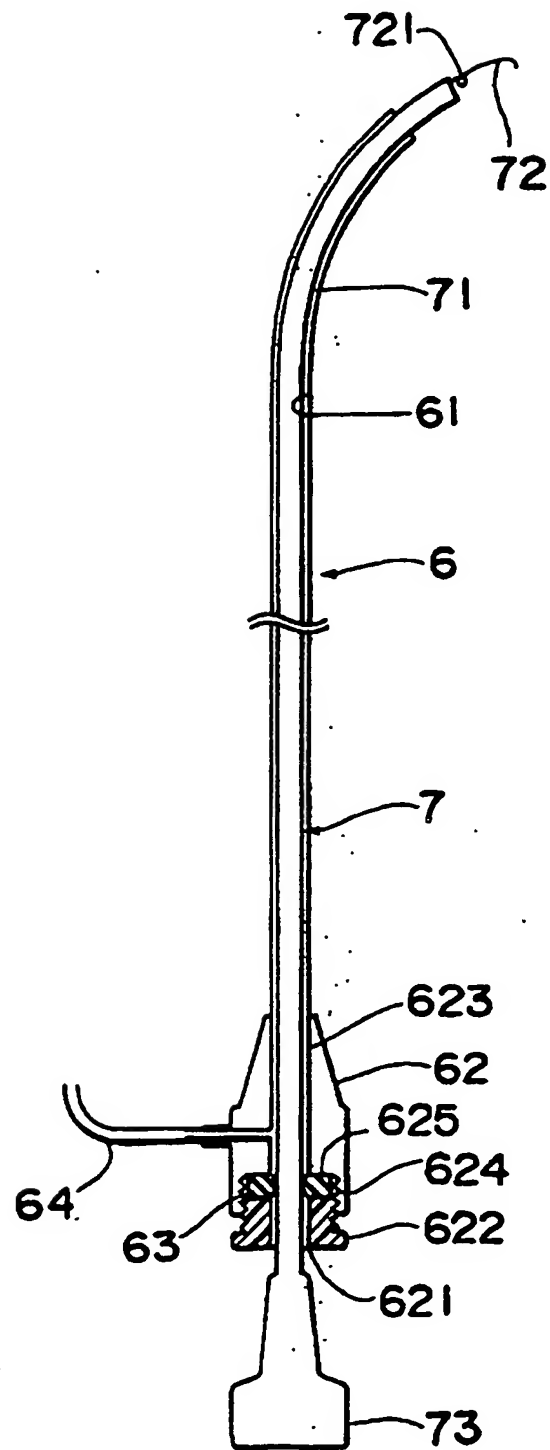


Fig. 5

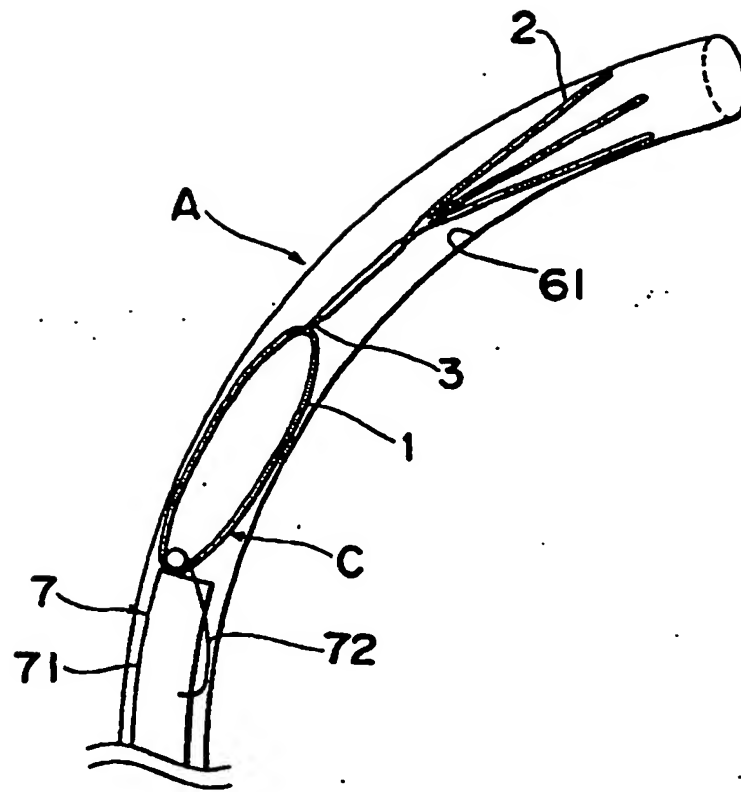


Fig. 6

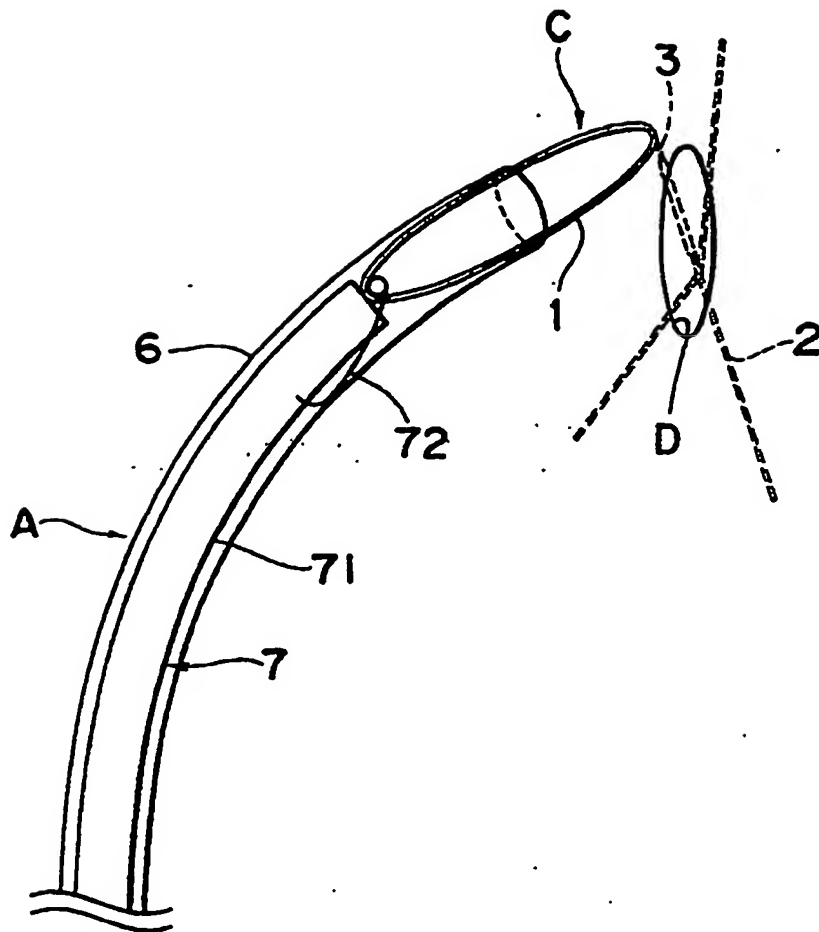


Fig. 7

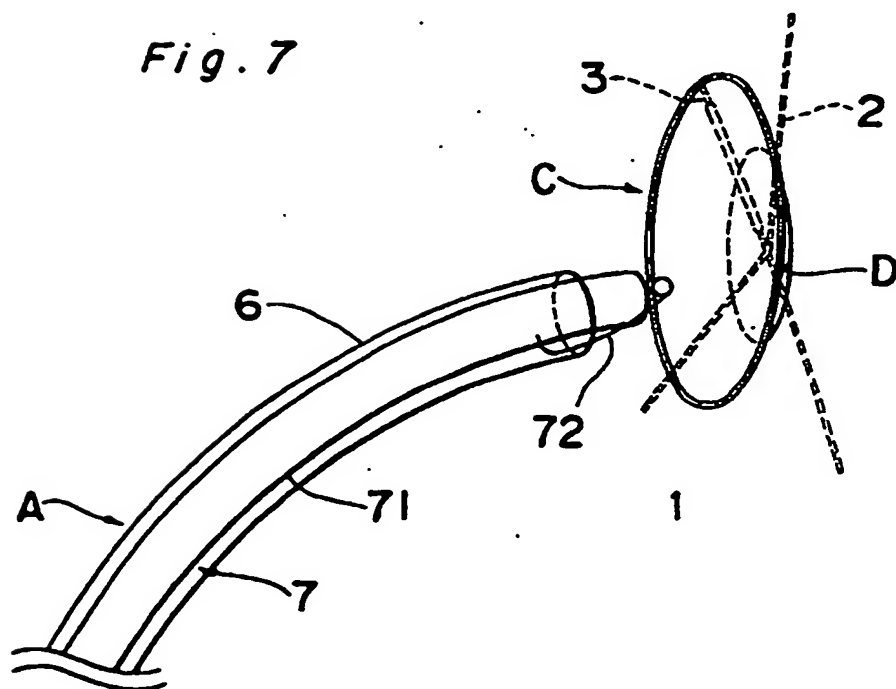


Fig. 8

